

Acanoides gen. n., a new spider genus from China with a note on the taxonomic status of *Acanthoneta* Eskov & Marusik, 1992 (Araneae, Linyphiidae, Micronetinae)

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Abstract

A new “micronetine” genus *Acanoides* gen. n. is erected to accommodate two species from China: *Acanoides beijingensis* sp. n. as the type species and *Acanoides hengshanensis* (Chen & Yin, 2000), comb. n., with the females described for the first time. The genitalic characters and somatic features of the new genus were studied by means of light microscopy and scanning electron microscopy (SEM). The monophyly of the new genus was tested by a phylogenetic analysis based on molecular data. Descriptions of the new genus, the new species and the new combination are presented; SEM images and microscopy pictures of somatic and genitalic characters are provided in detail. To distinguish from other genera with similar genitalic characters, we compare the new genus with the species of *Acanthoneta* Eskov & Marusik, 1992, *Epibellowia* Tanasevitch, 1996 and *Wubanoides* Eskov, 1986. Four putative synapomorphies for *Acanoides* gen. n. are suggested to support its monophyly that could be tested in the future. Furthermore, redescriptions of the epigynal morphology of *Acanthoneta aggressa* Chamberlin & Ivie, 1943 (Nearctic) and on the male of *A. dokutchaevi* Eskov & Marusik, 1993 (Far East Asia, firstly recorded from China) are provided. Based on comparison with *Poeciloneta*, from which *Acanthoneta* stat. n. was separated by Saaristo and Tanasevitch (1996), a revised diagnosis is proposed to support the generic status.

Keywords

Taxonomy, new species, new genus, genitalic morphology, movable epigynum

Introduction

Micronetinae Hull, 1920 is a fairly large subfamily of Linyphiidae Blackwall, 1859, including 1199 species placed in 90 genera (Tanasevitch 2014). It was redelimited by Saaristo and Tanasevitch (1996), who established eight new genera for 89 species, and raised three subgenera to generic status. Subsequently, a series of revisions were published (e.g. Saaristo and Tanasevitch 2002a, 2002b; Saaristo and Marusik 2004; Saaristo et al. 2006; Tu et al. 2006; Tu and Li 2006) that resulted in many new genera and a new subfamily Ipinae Saaristo, 2007. Results of these series of review works have not been tested in a phylogenetic context; neither Micronetinae nor Ipinae monophyly, as well as that of the genera included in the two subfamilies have been tested so far.

Poeciloneta hengshanensis (Chen & Yin, 2000) from China, originally placed in *Lepthyphantes* Menge, 1866, has its male palp very similar to that of *Poeciloneta (Acanthoneta) aggressa* (Chamberlin & Ivie, 1943). *Acanthoneta* Eskov & Marusik, 1992 is one of the three genera raised from subgeneric status by Saaristo and Tanasevitch (1996) with the type species *Acanthoneta aggressa*. Tu et al. (2006) transferred *P. hengshanensis* to *Acanthoneta* based on the similarity of the male palpal morphology. However, raising *Acanthoneta* to a generic status “was not accompanied by a diagnosis or justification”, and hence not accepted in The World Spider Catalog (Platnick 2014). All members of *Acanthoneta* are currently placed in *Poeciloneta* Kulczyński, 1894.

Females of *P. hengshanensis* (previously unknown) were found in new material from China. However, its epigynal conformation is neither congruent with that of *P. aggressa*, nor with any other species of *Poeciloneta*. Based on the presence of an extensible basal part, the movable epigynum accords with the diagnosis of the subfamily Ipinae Saaristo, 2007 (for example *Ipa* Saaristo, 2007 and *Solenysa* Simon, 1894). Additionally, we found another new species with genitalic morphology very similar to that of *P. hengshanensis*: the male palpal morphology similar to *Acanthoneta* and a movable epigynum in accordance with ipaine type.

A new genus *Acanoides* gen. n. is erected here for these two species. To test the placement of the new genus within Linyphiidae, a phylogenetic analysis based on newly sequenced molecular data of the two species and that of other linyphiids downloaded from NCBI was conducted. In the present study, the two species and the new genus are described. Characters of copulatory organs and somatic features of both species are illustrated by means of SEM and light microscopy. To distinguish the new genus from other “micronetine” genera with similar male palpal morphology and ipaine genera with a similar movable epigynum, the new genus is compared with the genera *Acanthoneta* (Micronetinae), *Wubanoides* Eskov, 1986 and *Epibellowia* Tanasevitch, 1996 (Ipinae). Due to limited material available for examination, comparisons are largely based on descrip-

tions and illustrations provided by Tanasevitch (1996), Saaristo and Tanasevitch (2000) and Saaristo (2007); images of the epigynum of *Acanthoneta aggressa* and the male of *A. dokutchaevi* Eskov and Marusik, 1994 are presented here. Four putative synapomorphies are suggested for *Acanoides* gen. n. that could be tested in future study. In addition, diagnoses for *Acanthoneta* stat. n. are provided based on comparison with illustrations of genitalic characters provided by Saaristo and Tanasevitch (2000), to support its generic status proposed by Saaristo and Tanasevitch (1996). The composition and monophyly of both *Acanoides* gen. n. and *Acanthoneta* stat. n. could be tested in future study.

Materials and methods

Specimens were examined and measured using a Leica M205A stereomicroscope. Male palps and epigyna were examined after they were dissected from the body. Left structures (e.g. palps, legs, etc.) were depicted. Embolic divisions were excised by breaking the membranous column which connects the suprategulum and radix. Male palps and epigyna were cleared in methyl salicylate. Digital images were taken with a Leica DFC 500 camera, as composites of multiple focus images assembled using the software package LEICA APPLICATION SUITE. Scanning electron microscopy (SEM) images were taken using a S-3400N scanning electron microscope at the China Agricultural University. For SEM examination the specimens were prepared following Álvarez-Padilla and Hormiga (2008). SEM images of the embolic division taken from the right palp were mirrored to match those taken from the left palp. All measurements were taken with a micrometer and are expressed in millimeters. The leg measurements are given in the following sequence: total (femur, patella+tibia, metatarsus, tarsus). All specimens examined here are deposited in the College of Life Sciences, Capital Normal University, China (CNU) and in the College of Life Sciences, Hunan Normal University, China (HNU), except for the female material of *A. aggressa*, the epigynal pictures of which were provided by Don Buckle (Saskatoon, Canada). Distribution data for these species within China are presented at the provincial level. Terminology for the epigynal characters follows Tu and Hormiga (2010) and male palpal and somatic characters follows that of Saaristo and Tanasevitch (1996) and Hormiga (2000). Anatomical abbreviations used in the text and figures:

Somatic morphology

AER	anterior eye row
ALE	anterior lateral eye(s)
AME	anterior median eye(s)
AMED	diameter of AME
PER	posterior eye row
PLE	posterior lateral eye(s)

Male palp

AX	apex of embolus
DM	distal membrane of terminal apophysis
DSA	distal suprategular apophysis
EM	embolic membrane
EP	embolus proper
FiG	Fickert's gland
LC	lamella characteristic
P	paracymbium
PCA	proximal cymbial apophysis
PH	pit hook
R	radix
SE	serrated area on embolus
SPT	suprategulum
TA	terminal apophysis
TH	thumb of embolus

Epigynum

CO	copulatory opening
CG	copulatory groove
DP	dorsal plate
EA	extensible area of epigynal basal part
EB	epigynal basal part
FG	fertilization groove
MP	median plate
S	spermathecae
SC	scape
ST	stretcher
VP	ventral plate

Phylogenetic analysis

Based on the dataset of Arnedo et al. (2009) which includes 34 linyphiid taxa (*Erigone dentipalpis* was not included as it has only one of the five genes available), newly sequenced data of the two *Acanoides* and data of another 65 linyphiid taxa downloaded from NCBI were added. A total of 111 taxa were sampled in our matrix, ten outgroup taxa of other araneoid families as in that of Arnedo et al. (2009) and 101 ingroup taxa, which cover the representatives of all the seven subfamilies currently proposed; one *Solenysa*, as a representative of ipaine, and *Acanthoneta* were included to test the placement of *Acanoides*.

Five genes: cytochrome c oxidase subunit I (CO1) and 16S rRNA (16S) and three nuclear genes 18S rRNA (18S), 28S rRNA (28S) and Histone H3 (H3) were sequenced for *Acanoides beijingensis* sp. n. and *A. hengshanensis*. Molecular procedures for sequencing follow that of Arnedo et al. (2009), with the same molecular markers to maximize the overlap of dataset. Taxa sampled and sequence accession numbers are presented in Appendix - Table S1. Data were automatic multiple aligned using the computer program Clustal X version 1.81 (Thompson et al. 1997). Gaps were treated as missing data. Maximum Likelihood analysis was performed using RAxML v7.2.7 as implemented on the Cipres Gateway (Miller et al. 2010). Bootstrap support analysis was performed with the commands: raxmlHPC-HYBRID-7.3.1 -T 6 -s infile -n result -x 12345 -p 876 -f a -N 1000 -m GTRCAT -q part.

Results

All five genes were sequenced for *Acanoides beijingensis* sp. n. and *A. hengshanensis* (Appendix - Table S1). The monophyly of Linyphiidae and its sister relationship with Pimoidae were not recovered in the result of phylogenetic analysis as two outgroup taxa: cyatholipid *Alaranea* and theridiosomatid *Theridiosoma* are embedded within Linyphiidae (Appendix - Fig. S1). Besides some weakly supported deeper branches, four robustly supported clades are recognized: *Stemonyphantes* clade (clade S), “micronetines-erigonines” clade (clade ME), “linyphiines”-1 clade (clade L1) and “linyphiines”-2 (clade L2). For the seven subfamilies currently proposed, only Stemonyphantinae Wunderlich, 1986 (the *Stemonyphantes* clade) and Mynogleninae Lehtinen, 1967 are monophyletic, while the mynoglenines clade and the *Dubiaranea* clade fall into clades L1 and L2 respectively that make Linyphiinae Blackwall, 1859 become a paraphyletic group; taxa of Micronetinae form a paraphyletic group, nested with taxa of Ipinae and Erigoninae within clade ME. The two *Acanoides* species form a robustly supported monophyly, distantly related to *Acanthoneta* and *Solenysa*.

Discussion

The result of the phylogenetic analysis based on molecular data suggests that the new species from Beijing is the sister taxon of *P. hengshanensis* which had ever been transferred to *Acanthoneta* by Tu et al. (2006). The lineage comprised by the two species is distantly related to *Acanthoneta* sp. (Appendix - Fig. S1). Accordingly, we erected here *Acanoides* gen. n. to accommodate the two species: *A. beijingensis* sp. n. and *A. hengshanensis* comb. n. The three known *Acanthoneta* species have very distinct male palpal morphology, only differ from that of the type species in small details (Eskov and Marusik 1992, 1993). Regardless the *Acanthoneta* taxon is congeneric with, or is the type species *Acanthoneta aggresus*, the new genus differs from all the three known species of *Acanthoneta* as well as all other “micronetines” in the females having a movable

epigynum (Figs 4G, 5G) and the males having a longer and sharper embolus proper (Figs 2D, 3D) which generally is pointed in “micronetines” (Fig. 6F); Fickert’s gland located in the membranous area outside the radix (Figs 2D, 3D), rather than embedded within the radix as usually the case in “micronetines” (Fig. 6G). This suggests that the two species are not congeneric with *Acanthoneta*.

Our results suggest an unknown *Leptophantes* species as a sister group to the *Acanoides* clade. *Leptophantes* Menge, 1866, which includes almost 500 species, is not a natural group (Saaristo and Tanasevitch 1996). All *Leptophantes* species, except five, have been transferred or are waiting to be transferred to other genera (e.g. Saaristo and Tanasevitch 1996, 2002a, b; Saaristo and Marusik 2004; Tu et al. 2006). The position of *Leptophantes* sp. on the tree indicates it is neither *Acanthoneta*, nor *Leptophantes*. Nevertheless, without morphological data, we fail to determine whether *Leptophantes* sp. is as a sister group of, or a number of *Acanoides* gen. n., so the close relative of *Acanoides* remains unresolved.

The genitalic characters of *Acanoides* make its subfamily placement problematic due to the epigynal character in accordance with Ipinae type, but the male palpal morphology of the “micronetine” type. Redelimitation of Mironetinae (Saaristo and Tanasevitch 1996) and the series of revisions of “micronetine” genera (e.g. Saaristo and Tanasevitch 2002a, 2002b; Saaristo and Marusik 2004; Saaristo et al. 2006; Tu et al. 2006; Tu and Li 2006) resulted in many new genera and even a new subfamily Ipinae (Saaristo 2007). However, none of them has been tested in a phylogenetic framework. Results of the first phylogenetic analysis for linyphiids based on molecular data indicate that neither Mironetinae nor Ipinae is a monophyletic group (Arnedo et al. 2009). Such a result was recovered in the present study too: “micronetine” taxa formed a paraphyletic group, and movable epigynum independently evolved in *Acanoides* and *Solenysa* (Appendix - Fig. S1). The extensible solenoid serving as a synapomorphy for *Solenysa* (Tu & Hormiga, 2011), the ventrally folded extensible epigynal basal part, together with long and sharp embolus proper, slender and unbranched lamella characteristic, and outside radix located Fickert’s gland are four putative synapomorphies for *Acanoides* gen. n.

With greatly increased ingroup sampling, the result of the present study produce a similar topology with that of Arnedo et al. (2009): four strongly supported clades S, L1, L2 and ME that correspond to the *Stemonyphantes* clade, clades C and D, and the “micronetines-erigonines” clade in the latter (Appendix - Fig. S1). Most newly added taxa fell into the clade ME that enriched the topology. However, the problems left by the previous study (Arnedo et al. 2009), such as the monophyly of Linyphiidae, placements of the weakly supported deeper branches, and taxa of different subfamilies placed together rendering most of the traditionally recognized subfamilies non monophyletic, persist. Six of the seven subfamilies currently proposed are not monophyletic groups. The higher level relationships within linyphiids reflected by phylogenetic result are still far away from the classic subfamily system (see Millidge 1984, 1993; Saaristo and Tanasevitch 1996; Saaristo 2007). Nevertheless, revising the whole higher level linyphiid systematics is beyond the scope of the present study. In the text bellow we keep using Mironetinae and Ipinae following the current taxonomic system.

Although with ingroup sampling about two times increased, the sampling size of the current dataset seems not to be enough to resolve the placements of *Acanoides* and *Acanthoneta*, as well as *Poeciloneta*, from which *Acanthoneta* were separated (Eskov and Marusik 1992), their close relatives, and the relationships among them. To better understand the higher level phylogenetic relationships of linyphiid spiders, more information, such as morphology and behavior, and a comprehensive sampling design are necessary. Here, we provide four putative synapomorphies for the new genus *Acanoides* that could be tested in future phylogenetic studies.

Taxonomy

Linyphiidae Blackwall, 1859

Acanoides gen. n.

<http://zoobank.org/4632240B-5228-4EB7-A1BC-CBD9176FEC2B>

<http://species-id.net/wiki/Acanoides>

Type species. *Acanoides beijingensis* sp. n.

Composition. Two species, *Acanoides beijingensis* sp. n. and *Acanoides hengshanensis* (Chen & Yin, 2000) comb. n.

Diagnosis. The males of *Acanoides* gen. n. can be distinct from *Acanthoneta* by the sharp embolus proper, the slender lamella characteristica unbranched, and by the Fickert's gland located in the membranous area outside the radix (Figs 2D, 3D). The females can be distinguished by the ventrally folded extensible epigynal basal part (Figs 2F, 3F).

Description. Male total length 2.34–2.73; female total length 2.10–2.42. Carapace yellowish-brown. Male carapace unmodified. AMEs smallest, others subequal; from the dorsal view AER recurved, PER straight, eyes separated by AMEd, ALE and PLE juxtaposed. Chelicerae medium-sized, with strong stridulatory ridges, female fang groove with three promarginal and three retromarginal teeth in *A. beijingensis* sp. n., and two promarginal and two retromarginal teeth in *A. hengshanensis*. Chaetotaxy: Ti I–IV: 2-2-2-2; Mt I–IV: 1-1-1-1; Mt I of males with two rows of ventral bristles, one prolateral, one retrolateral (Fig. 1C, 1D); Tm I about 0.25, Tm IV absent. Both species have a haplotracheate system.

Male palp (Figs 2A–E, 3A–E, 4A–B, 5A–B). Cymbium with proximal apophysis. Paracymbium medium to large-sized, with one tooth on lateral margin. Distal suprategular apophysis not modified as pit hook, or absent. Embolic division: radix long and narrow, Fickert's gland located in the membranous area connecting radix and embolus; embolus wide and strongly sclerotized with serrated area, embolus proper sharp with a thumb and an apex at each side; lamella characteristica unbranched, long and narrow with sharp sclerotized apex, almost parallel to radix; terminal apophysis with distal membrane.

Epigynum (Figs 2F–H, 3F–G, 4G–H, 5G–H). Protruding, with deeply wrinkled basal part, extensible and ventrally folded in constricted state. Epigynum well sce-

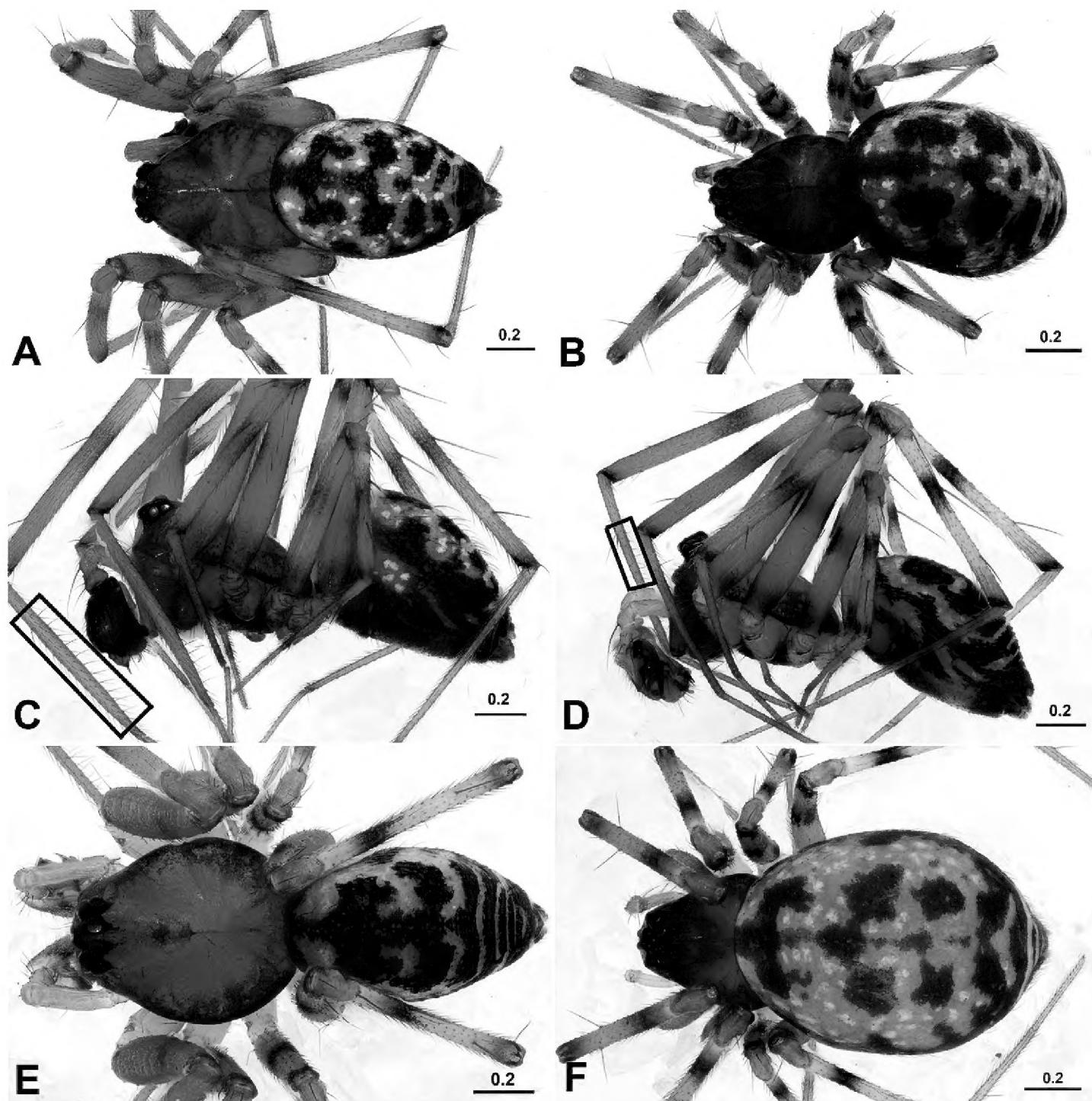


Figure 1. *Acanoides beijingensis* sp. n. (A–C) and *Acanoides hengshanensis* (D–F). **A** male, dorsal **B** female, dorsal **C** male, lateral, rectangle indicates ventrolateral rows of bristles on Mt I **D** male, lateral, rectangle indicates ventrolateral rows of bristles on Mt I **E** male, dorsal **F** female, dorsal. [Scale bars: mm].

rotized, epigynal cavity present (in *A. beijingensis* sp. n.) or absent (in *A. hengshanensis*), both scape and stretcher absent.

Etymology. The genus name, *Acanoides*, is a combination of the first four letters of “*Acanthoneta*” and the last five letters of “Wubanoides”. “-oides” itself in Latin means “similar to”, masculine in gender.

Phylogenetics. Due to limitations of the current dataset the monophyly of *Acanoides* could not be tested explicitly in our phylogenetic analyses, however it is supported by the following four putative synapomorphies: sharp embolus proper, slender and unbranched lamella characteristic, outside radix located Fickert’s gland and ventrally folded extensible epigynal basal part.

Distribution. China (Beijing, Hunan, Hebei) (Fig. 7).

Remarks. The males of *Acanoides* gen. n. have the palp of a “micronetine” type: presence of the Fickert’s gland, the boat-shaped radix, the trunk-like embolus with a pointed proper and a thumb, as well as the well developed terminal apophysis and lamella characteristica (Saaristo and Tanasevitch 1996). However, these sclerites in *Acanoides* (Fig. 2D) have some features different from the normal “micronetine” type (Fig. 6F, Saaristo and Tanasevitch 1996): Fickert’s gland is not embedded within the radix, but located in the membranous area connecting the radix and the embolus; and the embolus has a wide, strongly sclerotized body, with a longer and sharper embolus proper, whereas in most “micronetines” the embolus is usually trunk-like with a pointed embolus proper. The male palp of both *Acanoides* and *Acanthoneta*, have a long and slender lamella characteristica parallel to the long radix, but with an additional long and thin branch in *Acanthoneta* (Fig. 6F), unbranched in *Acanoides* (Figs 2D, 3D). The epigynum of *Acanthoneta* is in a normal “micronetine” type, with a sigmoid scape surrounded by an epigynal cavity (Fig. 6H), but with an extensible basal part in *Acanoides*.

The result of phylogenetic analysis based on molecular data indicates that Ipainae is not a monophyletic group as the movable epigynum independently evolved in *Acanoides* and *Solenysa* (Appendix - Fig. S1). This is also supported by the tracheal characters: haplotracheate type in *Acanoides*, but intermediate type in *Solenysa*, with the median pair extending into the prosoma (Tu and Hormiga 2011). We infer that the extensible basal part of the epigynum may have also evolved convergently with that in other ipaines. In *Acanoides* it differs by being ventrally folded, while it forms a solenoid in *Solenysa* (Tu & Hormiga, 2011), and folds inwards in other ipaines, e.g *Ipa* (Saaristo 2007: fig. 29), *Wubanoides* and *Epibellowia* (Tanasevitch 1996: figs 7–9). Furthermore, the male palp of typical ipaines has filiform embolus proper (Saaristo 2007: fig. 7; Tanasevitch 1996: figs 1, 4) much longer than that of *Acanoides* (Fig. 2D).

Acanoides beijingensis sp. n.

<http://zoobank.org/CE596A12-9C21-4B8F-97FC-F31CBC61CD7E>

http://species-id.net/wiki/Acanoides_beijingensis

Figs 1A–C, 2, 4

Type-locality. China, Beijing: Mt. Yangtaishan, 39°20.15'N; 115°34.52'E, alt. ca 320m, 15 Oct. 2007, L. Tu leg.

Type-specimens. Holotype, ♂ (CNU), China, Beijing, Mt. Yangtaishan, 39°20.15'N; 115°34.52'E, alt. ca 320 m, 15 Oct. 2007, L. Tu leg. Paratypes, 2 ♂♂ and 3 ♀♀ (CNU), same data as holotype.

Additional material examined. 1 ♂ and 2 ♀♀ (CNU), China, Hebei Province, Mt. Wulingshan, 40°33.61'N; 117°29.69'E, alt. ca 1100 m, 12 Aug. 2009, L. Tu leg.

Diagnosis. The male of *A. beijingensis* sp. n. can be distinguished from *A. hengshansensis* by the spine-shaped lamella characteristica (Figs 2D, 4C), ribbon-like in the latter (Figs 3D, 5C); by the hook-shaped terminal apophysis (Fig. 4C), straight in the latter (Fig. 5D); and by the presence of a distal suprategular apophysis (Fig. 4A), absent in

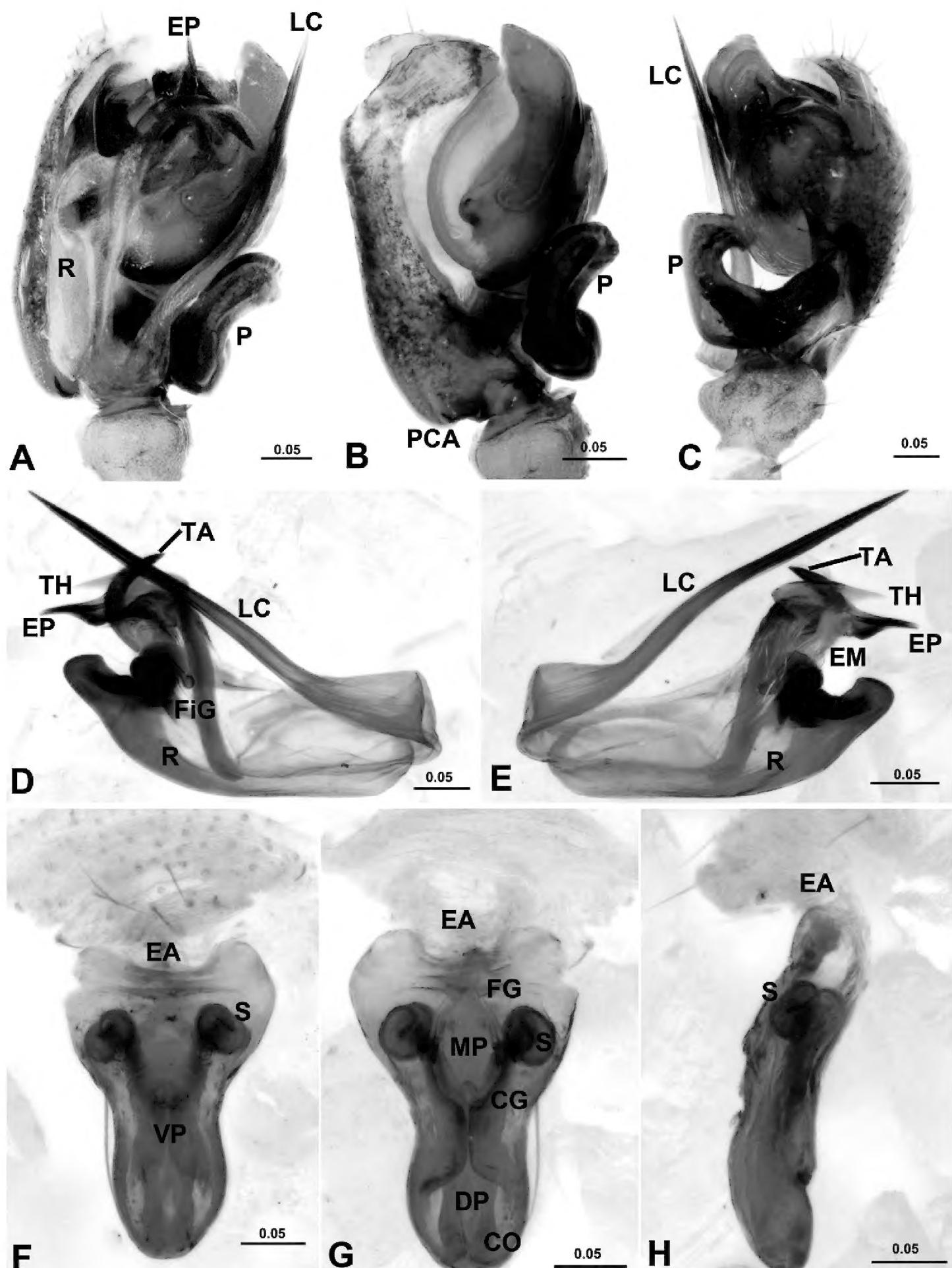


Figure 2. *Acanoides beijingensis* sp. n. **A** male palp, prolateral **B** male palp, prolateral, with embolic division removed **C** male palp, retrolateral **D** embolic division, ventral **E** embolic division, dorsal **F** epigynum, ventral **G** epigynum, dorsal **H** epigynum, lateral. CG copulatory groove; CO copulatory opening; DP dorsal plate; EA extensible area of epigynal basal part; EM embolic membrane; EP embolus proper; FG fertilization groove; FiG Fickert's gland; LC lamella characteristic; MP median plate; P paracymbium; PCA proximal cymbial apophysis; R radix; S spermathecae; TA terminal apophysis; TH thumb of embolus; VP ventral plate. [Scale bars: mm].

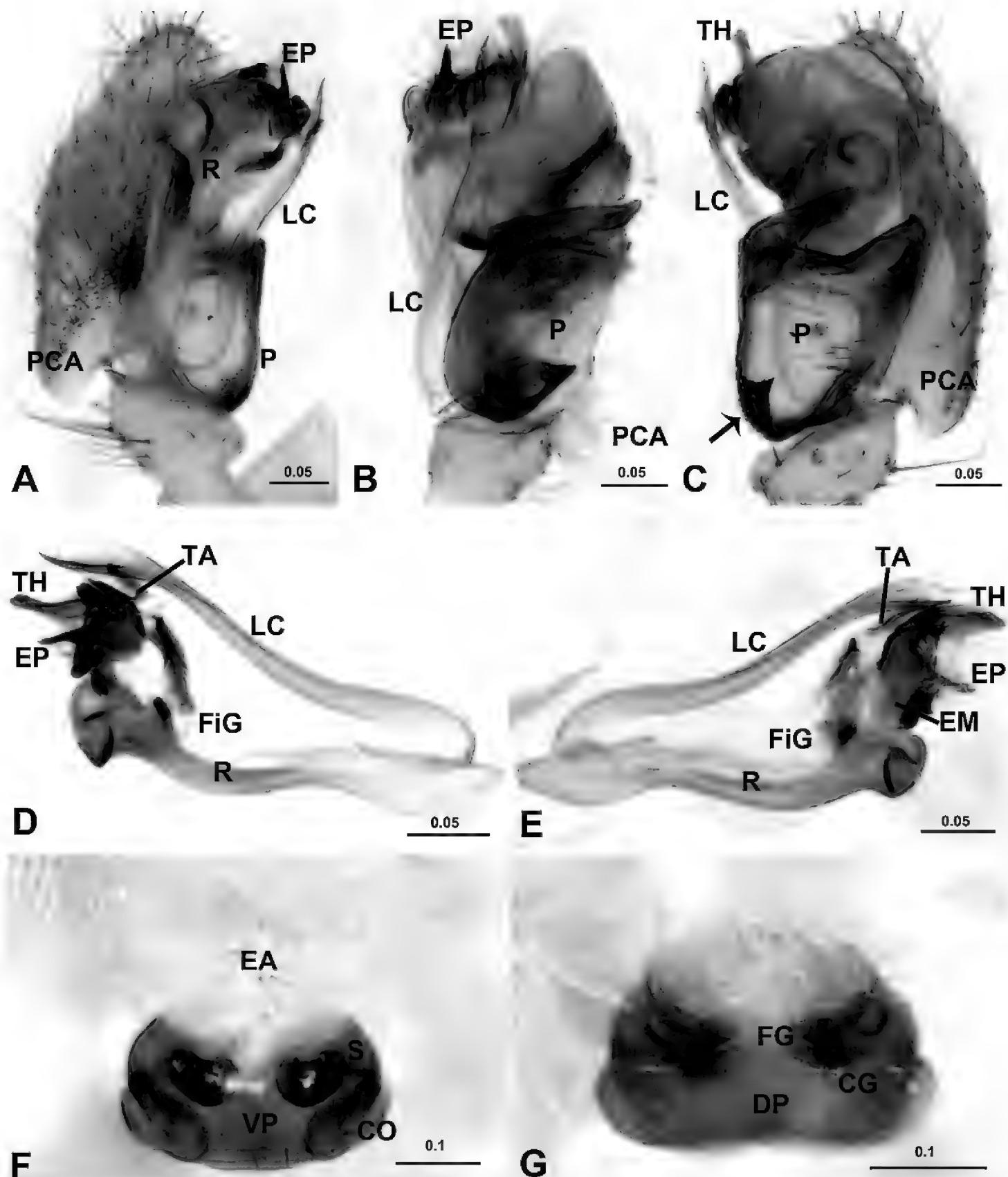


Figure 3. *Acanoides hengshanensis*. **A** male palp, prolateral **B** male palp, ventral **C** male palp, retrolateral, arrow indicates pointed tooth on posterolateral margin **D** embolic division, ventral **E** embolic division, dorsal **F** epigynum, ventral **G** epigynum, dorsal. CG copulatory groove; CO copulatory opening; DP dorsal plate; EA extensible area of epigynal basal part; EM embolic membrane; EP embolus proper; FG fertilization groove; FiG Fickert's gland; LC lamella characteristic; P paracymbium; PCA proximal cymbial apophysis; R radix; S spermatheca; TA terminal apophysis; TH thumb of embolus; VP ventral plate. [Scale bars: mm].

the latter. The female is distinct by having the epigynum two times longer than wide (Fig. 2F), shorter than wide in *A. hengshanensis* (Fig. 3F); and by the presence of a remnant epigynal cavity (Fig. 2G), totally absent in *A. hengshanensis* (Fig. 3G).

Description. Male holotype (Fig. 1A, C): Total length 2.69. Carapace 1.22 long, 1.01 wide. Abdomen 1.39 long, 0.88 wide. Lengths of legs: I 3.88 (1.05 + 1.18 + 0.99 + 0.66); II 3.02 (1.03 + 0.73 + 0.69 + 0.57); III 2.66 (0.87 + 0.88 + 0.51 + 0.40); IV 3.78 (1.12 + 1.09 + 0.93 + 0.64). Female (Fig. 1B): Total length 2.12. Carapace 0.93 long, 0.78 wide. Abdomen 1.25 long, 0.83 wide. Lengths of legs: I 6.10 (1.68 + 2.04 + 1.43 + 0.95); II 5.43 (1.56 + 1.74 + 1.24 + 0.89); III 4.39 (1.24 + 1.13 + 1.10 + 0.75); IV 5.88 (1.79 + 1.78 + 1.46 + 0.83). Tm I: 0.20. For other somatic features see description of the genus.

Male palp (Figs 2A–C, 4A–B). Cymbium with proximal apophysis. Paracymbium narrow, half rounded lateral tooth strongly sclerotized. Distal suprategular apophysis blunt, not modified as pit hook. Embolic division: radix long and narrow; Fickert's gland located in the membranous area connecting radix and embolus; embolus main body short and wide, strongly sclerotized, with serrated area on ventral surface; embolus proper sharp with pointed thumb and tail-like apex at each side; unbranched lamella characteristic long and slender, with sharp and strongly sclerotized apex; terminal apophysis hook-shaped with distal membrane.

Epigynum (Figs 2F–H, 4G–H). Two times longer than wide, wrinkled basal part extensible and ventrally folded in constricted state. Median plate and epigynal cavity present, without scape and stretcher. Copulatory openings opened dorsally.

Etymology. The species name refers to the type locality.

Variation. *Males* ($n = 3$). Total length 2.61–2.73. Carapace: 1.13–1.27 long, 0.95–1.05 wide. Abdomen 1.34–1.45 long, 0.71–0.99 wide.

Females ($n = 3$). Total length 2.10–2.23. Carapace: 0.90–0.96 long, 0.74–0.78 wide. Abdomen: 1.10–1.38 long, 0.79–0.88 wide.

Distribution. China (Beijing, Hebei) (Fig. 7).

Remarks. Although *A. beijingensis* sp. n. looks quite different from *A. hengshanensis* in the shape of the male paracymbium and in terms of female epigynal morphology, the strongly sclerotized embolus main body and the sharp embolus proper, the location of Fickert's gland, the presence of a ventrally folded extensive area of the epigynal basal part and the absence of a scape and stretcher, shared by the two species suggest they are closely related. A close relationship between the two species is additionally supported by the phylogenetic analysis (Appendix - Fig. S1).

Acanoides hengshanensis (Chen & Yin, 2000), comb. n.

http://species-id.net/wiki/Acanoides_hengshanensis

Figs 1D–F, 3, 5

Lepthyphantes hengshanensis Chen & Yin, 2000: 87, figs 12–16 (♂)

Acanthoneta hengshanensis: Tu et al. 2006: 412, figs 24–27 (♂).

Type-specimen. Holotype of *L. hengshanensis* Chen & Yin, 2000, ♂ (HNU), China, Hunan Province, Mt. Hengshan, 27°18'N; 112°42'E, 1–7 Aug. 1995, C. Yin leg. (examined).

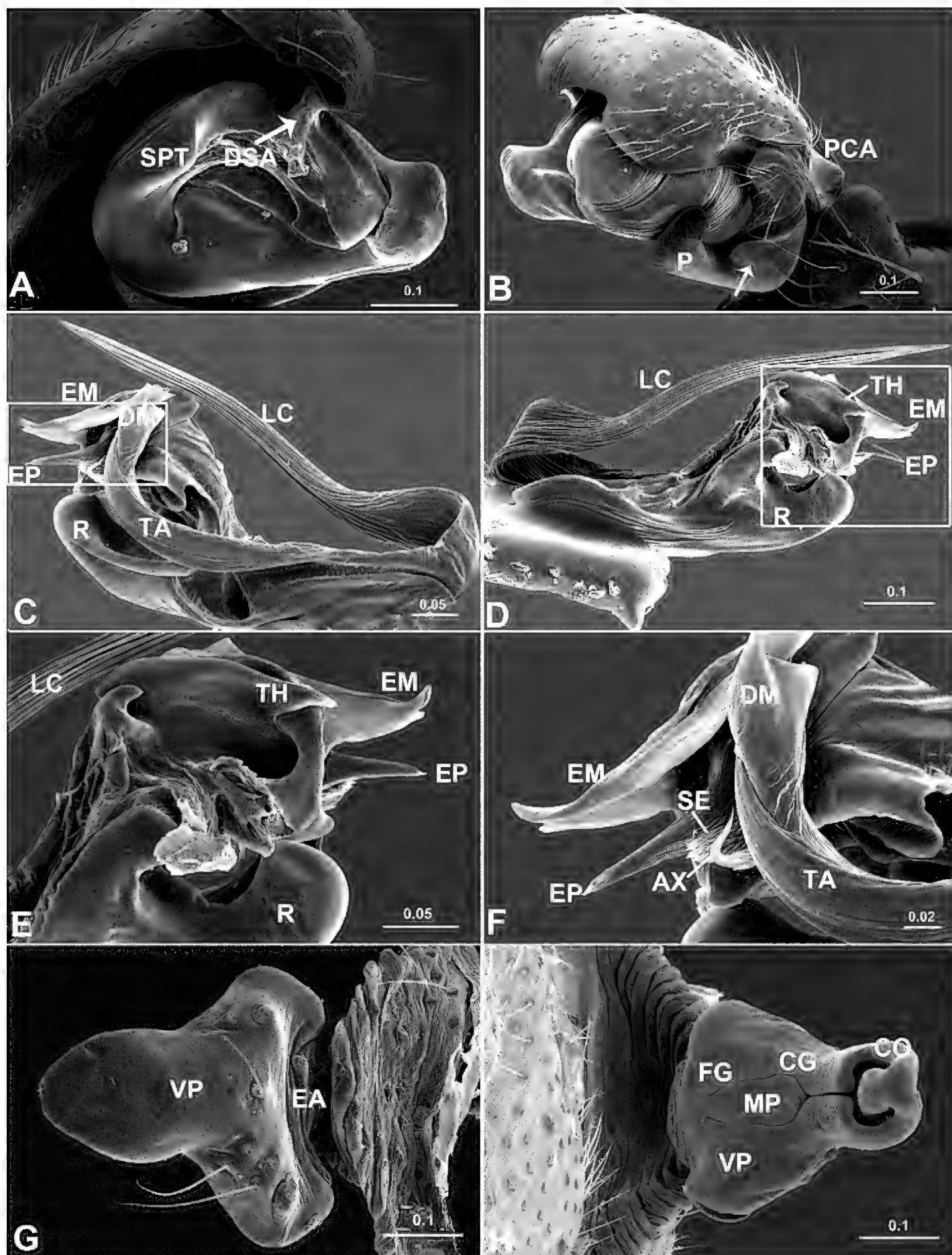


Figure 4. *Acanoides beijingensis* sp. n. **A** palp (embolic division removed), prolateral **B** palp, retrolateral, arrow indicates half rounded lateral tooth on paracymbium **C** embolic division, ventral **D** embolic division, dorsal **E** detail of **D** **F** detail of **C** **G** epigynum, ventral **H** epigynum, dorsal. AX apex of embolus; CG copulatory groove; CO copulatory opening; DM distal membrane of terminal apophysis; DSA distal suprategular apophysis; EA extensible area of epigynal basal part; EM embolic membrane; EP embolus proper; FG fertilization groove; LC lamella characteristic; MP median plate; P paracymbium; PCA proximal cymbial apophysis; R radix; S spermatheca; SE serrated area on embolus; SPT suprategulum; TA terminal apophysis; TH thumb of embolus; VP ventral plate. [Scale bars: mm].

Additional material examined. 3♂♂ and 4♀♀, China, Beijing, Mt. Yangtaishan, Dajue Temple, 40°03.06'N; 116°05.97'E, alt. 50 m, 15 Oct. 2007, L. Tu leg.

Diagnosis. See diagnosis for *A. beijingensis* sp. n.

Description. Male (Fig. 1D–E): Total length 2.39. Carapace 1.02 long, 0.78 wide. Abdomen 1.37 long, 0.78 wide. Lengths of legs: I 5.03 (1.37 + 1.56 + 1.32 + 0.78), II 3.33 (0.98 + 0.98 + 0.83 + 0.54), III 3.47 (0.98 + 1.07 + 0.88 + 0.54), IV 4.63 (1.27 + 1.41 + 1.22 + 0.73). Tm I: 0.24. Female (Fig. 1F): Total length 2.42. Carapace 0.96 long, 0.78 wide. Abdomen 1.80 long, 1.25 wide. Lengths of legs: I 4.21 (1.18 + 1.42 + 0.96 + 0.65), II 3.19 (0.98 + 1.06 + 0.66 + 0.49), III 2.81 (0.84 + 0.85 + 0.68 + 0.44), IV 3.70 (1.08 + 1.19 + 0.89 + 0.54). Tm I: 0.23. For other somatic characters see description of the genus.

Male palp (Figs 3A–C; 5A–B). Cymbium with distinct proximal apophysis pointing backwards. Paracymbium wide and U-shaped, with triangular tooth on postero-lateral margin. Distal suprategular apophysis absent. Embolic division: radix long and narrow; Fickert's gland located in the membranous area connecting radix and embolus; embolus main body large and strongly sclerotized with serrated area; embolus proper sharp with large thumb and pointed apex; lamella characteristic long and slender with bifurcated ends, one sharp and sclerotized, one membranous; terminal apophysis straight, with distal membrane.

Epigynum (Figs 3F–G, 5G–H). Short and wide, box-shaped, strongly sclerotized; wrinkled basal part extensible and ventrally folded in constricted state. Neither median plate nor epigynal cavity present. Copulatory openings located on ventral surface, slits of epigynal grooves extending laterally, passing from ventral to dorsal surface, then convergent mesally. No scape, no stretcher.

Variation. *Males* (n = 3). Total length 2.34–2.41. Carapace: 1.09–1.12 long, 0.72–0.93 wide. Abdomen 1.14–1.42 long, 0.68–0.83 wide.

Females (n = 4). Total length 2.32–2.42. Carapace: 0.87–1.01 long, 0.75–0.81 wide. Abdomen: 1.63–1.82 long, 0.76–1.22 wide.

Distribution. China (Beijing, Hunan) (Fig. 7).

Genus *Acanthoneta* Eskov & Marusik, 1992 stat. n.

Acanthoneta Eskov & Marusik, 1992: 34. Described as a subgenus of *Poeciloneta*.

Acanthoneta: Saaristo and Tanasevitch 1996: 175. Raised to generic status without any comments or argumentation.

Type species. *Poeciloneta aggressus* (Chamberlin & Ivie, 1943).

Composition. Three species: *A. aggressa* Chamberlin & Ivie, 1943 (Nearctic), *A. dokutchaevi* Eskov & Marusik, 1993 (Far East Asia) and *A. furcata* Emerton, 1913 (Nearctic).

Comments. Originally *Acanthoneta* was described as a subgenus of *Poeciloneta*, including two species: *Poeciloneta* (*A.*) *aggressa* and *Poeciloneta* (*A.*) *furcata*. One ad-

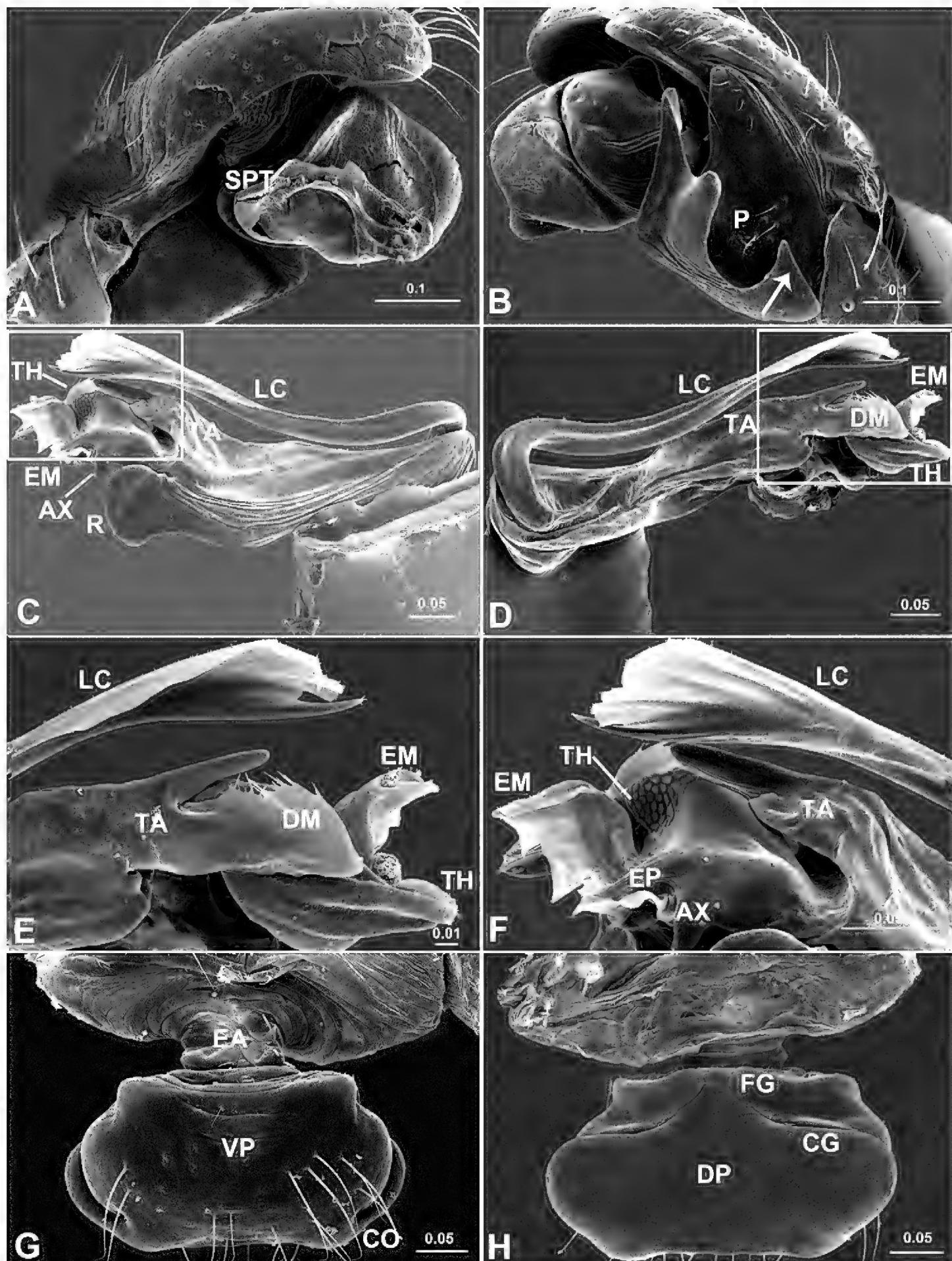


Figure 5. *Acanoides hengshanensis*. **A** palp (embolic division removed), prolateral **B** palp, retrolateral, arrow indicates pointed tooth on posterolateral margin **C** embolic division, ventral **D** embolic division, dorsal **E** detail of D **F** detail of C **G** epigynum, ventral **H** epigynum, dorsal. AX apex of embolus; CG copulatory groove; CO copulatory opening; DM distal membrane of terminal apophysis; EA extensible area of epigynal basal part; EM embolic membrane; EP embolus proper; FG fertilization groove; LC lamella characteristica; P paracymbium; PCA proximal cymbial apophysis; R radix; S spermatheca; SPT suprategulum; TA terminal apophysis; TH thumb of embolus; VP ventral plate. [Scale bars: mm].

ditional species *A. dokutchaevi* was assigned to the subgenus by Eskov and Marusik (1993). Saaristo and Tanasevitch (1996) raised *Acanthoneta* to genus status without any argumentations and hence the new status was not accepted by Platnick (2014). Here we provide a diagnosis for *Acanthoneta* and a comparison with *Poeciloneta*.

Diagnosis. Males of *Acanthoneta* differ from *Poeciloneta* by the long radix almost parallel with the long lamella characteristic (Fig. 6F), in the latter the radix is normal boat-shaped, lamella characteristic large and ribbon-like (Saaristo and Tanasevitch 2000: fig. 11). Females of the two genera differ by the epigynum in *Acanthoneta* having a sigmoid scape surrounded by an epigynal cavity, the lateral wall of which is posteriorly extended (Fig. 6H), whereas in *Poeciloneta* the scape is exposed, enlarged and strongly sclerotized (Saaristo and Tanasevitch 2000: fig. 18).

Acanthoneta aggressa (Chamberlin & Ivie, 1943)

http://species-id.net/wiki/Acanthoneta_aggressa

Fig. 6H–J

Lepthyphantes aggressus Chamberlin & Ivie, 1943: 14, figs 19–20.

Poeciloneta aggressa: Crawford 1988: 19.

Acanthoneta aggressa: Saaristo and Tanasevitch 1996: 175.

Poeciloneta aggressa: Paquin and Dupérré 2003: 147, figs 1623–1625.

Material examined. No material examined, epigynum pictures were provided by Don Buckle (Saskatoon, Canada): 1 ♀, Canada, Alberta, Chinook Lake, under log in spruce or fir woods, 49°40'N; 114°30'W, 25 Jul. 1988, D. J. Buckle leg.

Description. *Epigynum* (Fig. 6H–J). Slightly protruding, without extensible area at basal part. Epigynal cavity, with posteriorly extended lateral wall, surrounding sigmoid folded scape; scape long and narrow, with well developed lateral lobes hosting copulatory openings and distal stretcher.

Distribution. Across North America from Washington State to Québec (Buckle et al. 2001; Paquin and Dupérré 2003).

Acanthoneta dokutchaevi Eskov & Marusik, 1993

http://species-id.net/wiki/Acanthoneta_dokutchaevi

Fig. 6A–G

Poeciloneta (Acanthoneta) aggressa non Chamberlin & Ivie, 1943: Eskov and Marusik 1992: 34–35, figs 11–13 (♂).

Poeciloneta (Acanthoneta) dokutchaevi: Eskov and Marusik 1993: 52, figs 49–51 (♂).

Material examined. 1 ♂, China, Jilin Province, Mt. Changbaishan, Ski. 42°01.54'N; 128°04.25'E, alt. ca 1260 m, 31 July 1971.

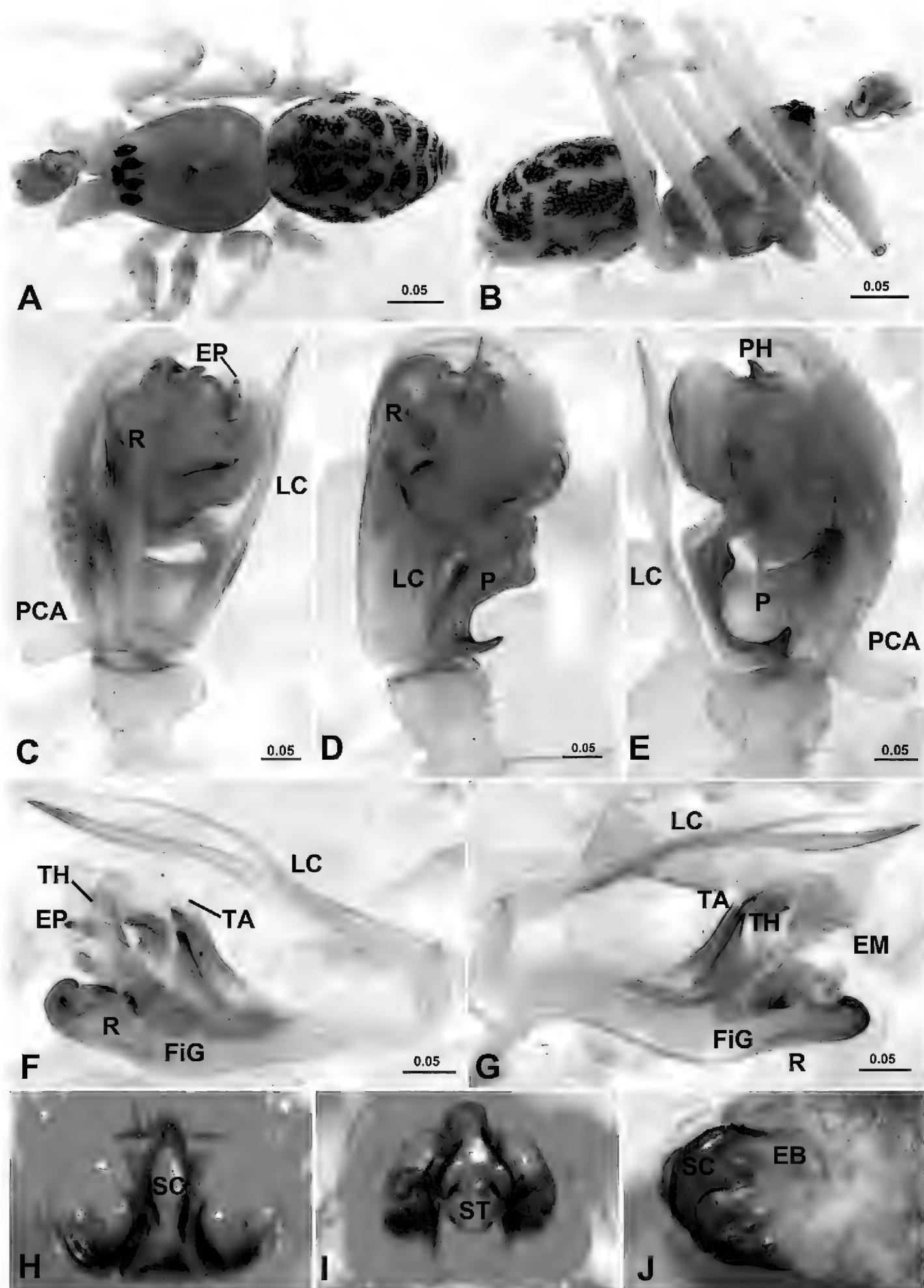


Figure 6. *Acanthoneta dokutchaevi* (A–G) and *Acanthoneta aggressa* (H–J). **A** male, dorsal **B** male, lateral **C** male palp, prolateral **D** male palp, ventral **E** male palp, retrolateral **F** embolic division, ventral **G** embolic division, dorsal **H** epigynum, ventral **I** epigynum, posterior **J** epigynum, lateral (**H–J** photos provided by Don Buckle). EB epigynal basal part; EM embolic membrane; EP embolus proper; FiG Fickert's gland; LC lamella characteristica; P paracymbium; PCA proximal cymbial apophysis; PH pit hook; R radix; SC scape; ST stretcher; TA terminal apophysis; TH thumb of embolus. [Scale bars: mm].

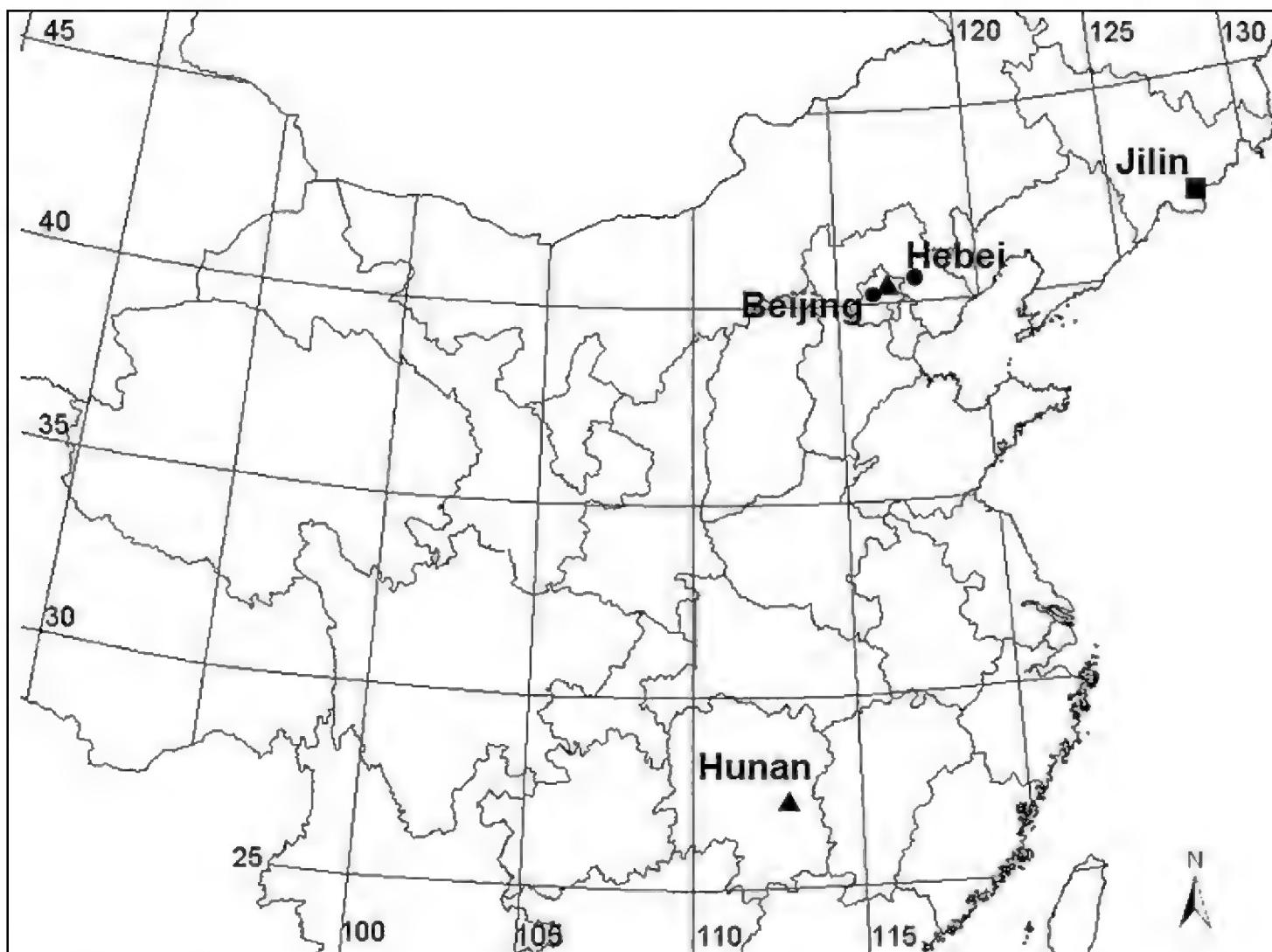


Figure 7. Collecting localities of *Acanoides* species and *Acanthoneta aggressa*, *Acanoides beijingensis* sp. n. (Beijing, Hebei); *Acanoides hengshanensis* (Hunan, Beijing); *Acanthoneta dokutchaevi* (Jilin).

Description. Male (Fig. 6A–B). Chelicera long, with strong stridulatory ridges. Chaetotaxy: Ti I–IV: 2-2-2-2; Mt I–IV: 1-0-0-1; Tm I about 0.80, Tm IV present. For other somatic characters see description by Eskov and Marusik (1993).

Male palp (Fig. 6C–E). Cymbium with proximal apophysis erected. Paracymbium wide, with two pointed teeth on lateral margin. Distal suprategular apophysis modified as pit hook. Embolic division: radix long and narrow; Fickert's gland located within radix; embolus main body trunk-like with serrated area, pointed embolus proper and well developed thumb; lamella characteristic fork-like branched, long and slender, almost parallel to radix; terminal apophysis with distal membrane and two strongly sclerotized teeth on ventral side.

Female. Unknown.

Remarks. The male of this species is similar to the type species *A. aggressa*. It differs only by the shape of the paracymbium. For a detailed comparison see Eskov and Marusik (1993).

Distribution. Far East Asia: Magadan Area (Eskov and Marusik 1993) and China (Fig. 7) (new record).

Acknowledgements

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References

- Álvarez-Padilla F, Hormiga G (2008) A protocol for digesting internal soft tissues and mounting spiders for scanning electron microscopy. *Journal of Arachnology* 35: 538–542. doi: 10.1636/Sh06-55.1
- Arnedo MA, Hormiga G, Scharff N (2009) Higher level phylogenetics of linyphiid spiders (Araneae, Linyphiidae) based on morphological and molecular evidence. *Cladistics* 25: 231–262. doi: 10.1111/j.1096-0031.2009.00249.x
- Buckle DJ, Carroll D, Crawford RL, Roth VD (2001) Linyphiidae and Pimoidae of America north of Mexico: checklist, synonymy, and literature. *Fabreries Supplement* 10: 89–191.
- Chamberlin RV, Ivie W (1943) New genera and species of North American Linyphiid spiders. *Bulletin of the University of Utah* 7: 1–39.
- Chen J, Yin C (2000) On five species of linyphiid spiders from Hunan, China (Araneae: Linyphiidae). *Acta Arachnologica Sinica* 9: 86–93.
- Edgar RC (2004) Muscle: multiple sequence alignment with high accuracy and high throughput. *Nucleic Acids Research* 32: 1792–1797. <http://www.ebi.ac.uk/Tools/muscle/> doi: 10.1093/nar/gkh340
- Eskov KY, Marusik YM (1993) New data on the taxonomy and faunistics of North Asian linyphiid spiders (Aranei Linyphiidae). *Arthropoda Selecta* 2: 41–79.
- Eskov KY, Marusik YM (1992) On the mainly Siberian spider genera *Wubanoides*, *Parawubanoides* gen. n. and *Poeciloneta*. *Arthropoda Selecta* 1: 21–38.
- Hormiga G (2000) Higher level phylogenetics of erigonine spiders (Araneae, Linyphiidae, Erigoninae). *Smithsonian Contributions to Zoology* 609: 1–160. doi: 10.5479/si.00810282.609
- Miller JR, Koren S, Sutton G (2010) Assembly algorithms for next-generation sequencing data. *Genomics* 95: 315–327. doi: 10.1016/j.ygeno.2010.03.001
- Millidge AF (1984) The taxonomy of the Linyphiidae, based chiefly on the epigynal and tracheal characters (Araneae: Linyphiidae). *Bulletin of British Arachnological Society* 6: 229–267.
- Millidge AF (1993) Further remarks on the taxonomy and relationships of the Linyphiidae, based on the epigynal duct conformation and other characters. *Bulletin of British Arachnological Society* 9: 145–156.

- Paquin P, Dupérré N (2003) Guide d'identification des Araignées (Araneae) du Québec. Fabreries Supplement 11: 147.
- Platnick NI (2014) The world spider catalog, version 13.0. American Museum of Natural History. <http://research.amnh.org/iz/spiders/catalog/LINYPHIIDAE.html>
- Saaristo MI (2007) A new subfamily of linyphiid spiders based on a new genus created for the keyserlingi-group of the genus *Lepthyphantes* (Aranei: Linyphiidae). Arthropoda Selecta 16: 33–42.
- Saaristo MI, Marusik YM (2004) Two new petrophilous “micronetine” genera, *Agypantes* gen. n. and *Lidia* gen. n. (Araneae, Linyphiidae, Micronetinae), from the eastern Palearctic with descriptions of two new species. Bulletin of British Arachnological Society 13: 76–82.
- Saaristo MI, Tu LH, Li SQ (2006) A review of Chinese micronetine species (Araneae: Linyphiidae). Part I: Species of ex-*Arcuphantes* and ex-*Centronerus*. Animal Biology 56: 383–401. doi: 10.1163/157075606778441886
- Saaristo MI, Tanasevitch AV (2002a) A new micronetid spider genus from the Oriental Region (Aranei: Linyphiidae: Micronetinae). Arthropoda selecta 11: 319–330.
- Saaristo MI, Tanasevitch AV (2002b) *Helsingenia* gen. n., a new micronetid genus from Old-World tropics (Aranei: Linyphiidae: Micronetinae). Arthropoda selecta 11: 153–158.
- Saaristo MI, Tanasevitch AV (2000) Systematics of the *Bolyphantes-Poeciloneta* genus group of the subfamily Micronetinae Hull, 1920 (Arachnida: Araneae: Linyphiidae). Reichenbachia 33: 255–265.
- Saaristo MI, Tanasevitch AV (1996) Redelimitation of the Subfamily Micronetinae Hull, 1920 and the Genus *Lepthyphantes* Menge, 1866 with Descriptions of Some New Genera (Aranei, Linyphiidae). Berichte naturw.-mediz. Verein Innsbruck 83: 163–186.
- Tanasevitch AV (2014) Linyphiid spiders of the world. <http://www.andtan.newmail.ru/list/>
- Tanasevitch AV (1996) Reassessment of the spider genus *Wubanoides* Eskov, 1986 (Arachnida: Araneae: Linyphiidae). Reichenbachia Mus. Tierkd. Dresden 1996: 123–129.
- Thompson JD, Gibson TJ, Plewniak F, Jeanmougin F, Higgins DG (1997) The CLUSTAL_X windows interface: flexible strategies for multiple sequence alignment aided by quality analysis tools. Nucleic Acids Research 24: 4876–4882. doi: 10.1093/nar/25.24.4876
- Tu LH, Hormiga G (2011) Phylogenetic analysis and revision of the linyphiid spider genus *Solenysa* (Araneae: Linyphiidae: Erigoninae). Zoological Journal of the Linnean Society 161: 484–530. doi: 10.1111/j.1096-3642.2010.00640.x
- Tu LH, Hormiga G (2010) The female genitalic morphology of “micronetine” spiders (Araneae, Linyphiidae). Genetica 138: 59–73. doi: 10.1007/s10709-009-9368-9
- Tu LH, Saaristo MI, Li SQ (2006) A review of Chinese micronetine species (Araneae: Linyphiidae). Part II: Seven species of ex-*Lepthyphantes*. Animal Biology 56: 403–421. doi: 10.1163/157075606778441877
- Tu LH, Li SQ (2006) A new *Drapetisca* species from China and comparison with European *D. socialis* (Sundevall, 1829) (Araneae: Linyphiidae). Animal Biology 56: 383–401.

Appendix

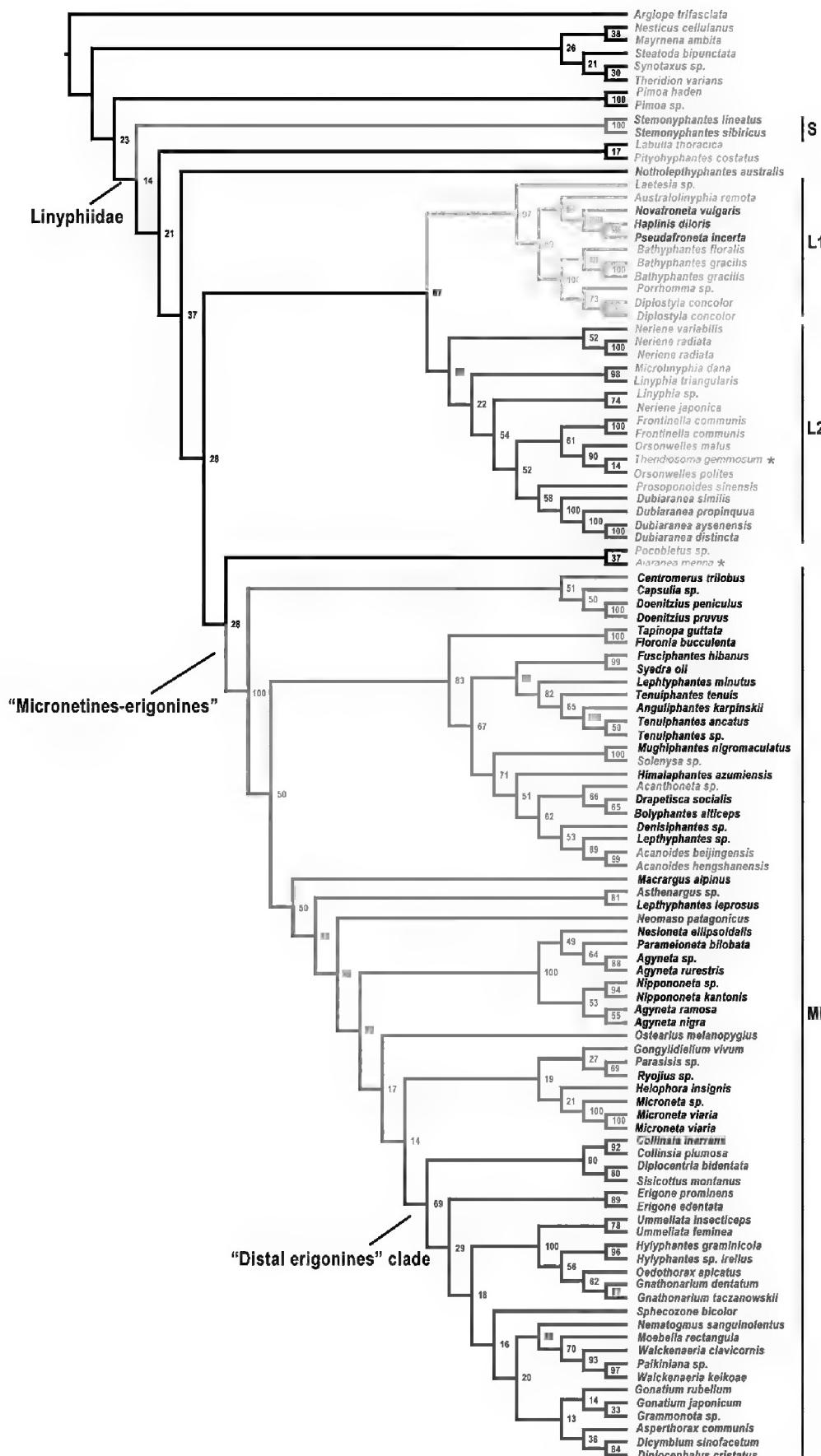


Figure S1. Linyphiid phylogeny resulting from Maximum Likelihood analysis based on molecular data. Numbers at the nodes are bootstrap value. Branches in color indicate the four robustly supported clades within linyphiids: **S** *Stemonyphantes* clade (blue) **L1** “linyphiines”-1 clade (pale green) **L2** “linyphiines”-2 clade (dark blue) **ME** “micronetines-erigonines” clade (red, with “Distal erigonines” clade in green). Taxa in different colors sampled from different groups: grey, outgroup; blue, Stemonyphantinae; pale green, Linyphiinae; dark blue, Mynogleninae; pink, Dubiaraneinae; black, Micronetinae; red, Ipinae, *Acanthoneta* and *Acanoides* gen. n.; green, Erigoninae. Red stars indicate the two out-group taxa: cyatholipid *Alaranea* and theridiosomatid *Theridiosoma* embedded within Linyphiidae.

Table S1. GenBank accession numbers. Data of the taxa labeled with “#” are newly sequenced; the taxa labeled with “**” come from Arnedo et al. 2009.

Family	Genus	Species	16s	18s	28s	COI	H3
Araneidae	<i>Argiope</i>	<i>trifasciata</i>		FJ525386		FJ525316	FJ525335
Cyatholipidae	<i>Alarnea</i>	<i>merina</i> *	AY230942	AY230890	AY231074	AY231022	AY230982
Mysmenidae	<i>Maymena</i>	<i>ambita</i>	GU456746	GU456765	GU456824	GU456876	GU456921
Nesticidae	<i>Nesticus</i>	<i>cellulanus</i>	EU746444	AF005447	AF124961	EU746435	
Pimoidae	<i>Pimoa</i>	<i>haden</i>	GU338640	GU338524	EF128112	EF128155	
Pimoidae	<i>Pimoa</i>	sp.*	AY230940	AY230893	AY231072	AY231025	AY230985
Synotaxidae	<i>Synotaxus</i>	sp.	AY230943	AY230894	AY231076	AY231026	AY230986
Theridiidae	<i>Steatoda</i>	<i>bipunctata</i> *	AY230951	AY230926	AY231084	AY231057	AY231014
Theridiidae	<i>Theridion</i>	<i>varians</i> *	AY230976	AY230932	AY231111	AY231063	AY231017
Theridiosomatidae	<i>Theridiosoma</i>	<i>gemmaosum</i>	HM030408	HM030417	HM030428	HM030436	HM030443
Linyphiidae	<i>Acanoides</i>	<i>beijingensis</i> #	KJ027589	KJ027587	KJ027580	KJ027582	KJ027583
	<i>Acanoides</i>	<i>hengshanensis</i> #	KJ027585	KJ027588	KJ027584	KJ027586	KJ027581
	<i>Acanthoneuta</i>	sp.		GU338479	GU338560	GU338678	
	<i>Agynteta</i>	sp.	GU338621		GU338529		
	<i>Agynteta</i>	<i>ramosa</i> *	FJ838670	FJ838694	FJ838717	FJ838648	FJ838740
	<i>Anguliphantes</i>	<i>karpinskii</i>		GU338516	GU338566	GU338680	
	<i>Asperthorax</i>	<i>communis</i>		GU338482	GU338545	GU338684	
	<i>Asthenargus</i>	sp.		GU338493	GU338561		
	<i>Australolinyphia</i>	<i>remota</i> *	FJ838671	FJ838695	FJ838718	FJ838649	FJ838741
	<i>Bathyphantes</i>	<i>floralis</i>	GU338604	GU338465	GU338583	GU338659	
	<i>Bathyphantes</i>	<i>gracilis</i> *	FJ838672	FJ838696	FJ838719	FJ838650	FJ838742
	<i>Bathyphantes</i>	<i>gracilis</i>	GU338630	GU338464	GU338582	GU338689	
	<i>Bathyphantes</i>	<i>alticeps</i> *	AY078660	AY078667	AY078678	AY078691	AY078700
	<i>Capulia</i>	sp.		GU338470	GU338586		
	<i>Centromerus</i>	<i>trilobus</i>	GU338599	GU338468	GU338571	GU338656	
	<i>Collinsia</i>	<i>inerrans</i>	GU338601	GU338518		GU338645	
	<i>Collinsia</i>	<i>plumose</i>	GU338638	GU338499	GU338543		
	<i>Denisiphantes</i>	sp.	GU338619	GU338508	GU338663	GU338669	
	<i>Dicymbium</i>	<i>sinfacatum</i>	GU338614	GU338487	EF128119	GU338665	

Family	Genus	Species	16s	18s	28s	COI	H3
	<i>Diplocentria</i>	<i>bidentata</i>	GU338629	GU338494	GU338542	GU338688	
	<i>Diplocephalus</i>	<i>cristatus</i>	GU338637	GU338490		GU338696	
	<i>Diplostyla</i>	<i>concolor*</i>	FJ838673	FJ838697	FJ838720	FJ838651	FJ838743
	<i>Diplostyla</i>	<i>concolor</i>	GU338639	GU338467	GU338585		GU338697
	<i>Doenitzius</i>	<i>peniculus</i>	GU338631	GU338469		GU338690	
	<i>Doenitzius</i>	<i>pruinus</i>	GU338632	GU338474		GU338691	
	<i>Drapetisca</i>	<i>socialis*</i>	FJ838674	FJ838698	FJ838721	FJ838652	FJ838744
	<i>Dubiaranea</i>	<i>aysenensis</i>	FJ838675	FJ838699	FJ838722	FJ838653	FJ838745
	<i>Dubiaranea</i>	<i>distincta</i>	GU338624	GU338459	GU338579		GU338648
	<i>Dubiaranea</i>	<i>propinquua</i>	GU338627	GU338460	GU338580		GU338675
	<i>Dubiaranea</i>	<i>similis</i>		GU338521	GU338581		GU338681
	<i>Erigone</i>	<i>edentata</i>		GU338486	GU338540		GU338686
	<i>Erigone</i>	<i>prominens</i>		GU338498	GU338539		GU338679
	<i>Floronia</i>	<i>bucculenta*</i>	FJ838676	FJ838700	FJ838723	FJ838654	FJ838746
	<i>Frontinella</i>	<i>communis</i>	GU338628	GU338517	GU338573		
	<i>Frontinella</i>	<i>communis*</i>	FJ838677	FJ838701	FJ838724	FJ838655	FJ838747
	<i>Fusciphantes</i>	<i>hibanus</i>		GU338512	GU338570		GU338683
	<i>Gnathonarium</i>	<i>dentatum</i>	GU338593	GU338477	EF128120		GU338651
	<i>Gnathonarium</i>	<i>taczanowskii</i>	GU338620	GU338480	GU338547		GU338670
	<i>Gonatium</i>	<i>japonicum</i>	GU338613	GU338492			
	<i>Gonatium</i>	<i>rubellum*</i>	FJ838679	FJ838703	FJ838726	FJ838656	FJ838749
	<i>Gongylidium</i>	<i>vivum*</i>	FJ838678	FJ838702	FJ838725		FJ838748
	<i>Grammonota</i>	sp.		GU338491		GU338685	
	<i>Haplinis</i>	<i>diloris*</i>	FJ838680	FJ838704	FJ838727	FJ838657	FJ838750
	<i>Helephora</i>	<i>insignis*</i>	FJ838681	FJ838705	FJ838728	FJ838658	FJ838751
	<i>Himalaphantes</i>	<i>azumiensis</i>		GU338522	GU338564		GU338677
	<i>Hylaphantes</i>	<i>graminicola</i>	GU338595	GU338478	GU338550		GU338653
	<i>Hylaphantes</i>	sp.	GU338618	GU338481	GU338549		GU338668
	<i>Labulla</i>	<i>thoracica*</i>	AY078662	AY078674	AY078680	AY078694	AY078707
	<i>Laetesia</i>	sp.*	FJ838682	FJ838706	FJ838729	FJ838659	FJ838752
	<i>Lepthyphantes</i>	<i>minutus*</i>	AY078663	AY078673	AY078681	AY078689	AY078705

Family	Genus	Species	16s	18s	28s	COI	H3
	<i>Lepthyphantes</i>	<i>leprosus</i>		GU338488	GU338565	GU338682	
	<i>Lepthyphantes</i>	sp.	GU338610	GU338509	GU338562	GU338664	
<i>Linyphia</i>		<i>triangularis</i> *	AY078664	AY078668	AY078682	AY078693	AY078702
<i>Linyphia</i>	sp.		GU338597	GU338461	GU338572	GU338654	
<i>Macrargus</i>	<i>alpinus</i>			GU338505	GU338559		
<i>Agyrta</i>	<i>nigra</i>	GU338608	GU338504	GU338577	GU338662		
<i>Agyrta</i>	<i>rurestris</i> *	FJ838683	FJ838707	FJ838730	FJ838660	FJ838753	
<i>Microlynphia</i>	<i>dana</i> *	AY078665	AY078677	AY078683	AY078690		
<i>Microneta</i>	sp.	GU338609	GU338472	GU338538	GU338663		
<i>Microneta</i>	<i>viaria</i> *	FJ838684	FJ838708	FJ838731	FJ838661	FJ838754	
<i>Microneta</i>	<i>viaria</i>	GU338598	GU338502	GU338537	GU338655		
<i>Moebelia</i>	<i>rectangular</i>	GU338591	GU338485	GU338557			
<i>Mugiphantes</i>	<i>nigromaculatus</i>	GU338600	GU338510	GU338527	GU338644		
<i>Nematognathus</i>	<i>sanguinolentus</i>	GU338635	GU338489	GU338544	GU338694		
<i>Neomaso</i>	<i>patagonicus</i>	GU338626	GU338473	GU338578			
<i>Neriene</i>	<i>japonica</i>	GU338633	GU338462	GU338575	GU338692		
<i>Neriene</i>	<i>radiata</i> *	AY078710	AY078670	AY078684	AY078696	AY078709	
<i>Neriene</i>	<i>radiate</i>	GU338623	GU338463	GU338574	GU338672		
<i>Neriene</i>	<i>variabilis</i> *	AY078711	AY078669	AY078685	AY078699	AY078706	
<i>Nesioneta</i>	<i>ellipsoidalis</i>		GU338519	GU338532	GU338687		
<i>Nippaponeta</i>	<i>kantonis</i>	GU338634	GU338471	GU338530	GU338693		
<i>Nippaponeta</i>	sp.	GU338602	GU338520	GU338531	GU338657		
<i>Notholephyphantes</i>	<i>australis</i> *	FJ838685	FJ838709	FJ838732	FJ838662	FJ838755	
<i>Novafroneta</i>	<i>vulgaris</i> *	FJ838686	FJ838710	FJ838733	FJ838663	FJ838756	
<i>Oedothorax</i>	<i>apicatus</i> *	FJ838687	FJ838711		FJ838664	FJ838757	
<i>Orsonwelles</i> *	<i>malus</i>	AY078737	AY078676	AY078795	AY078697	AY078708	
<i>Orsonwelles</i> *	<i>polites</i>	AY078725	AY078671	AY078786	AY078755	AY078701	
<i>Ostearius</i>	<i>melanopygius</i> *	FJ838688	FJ838712	FJ838735		FJ838758	
<i>Paikiniana</i>	sp.	GU338617	GU338495	GU338555	GU338647		
<i>Parameioneta</i>	<i>bilobata</i>	GU338605	GU338503	GU338533	GU338660		
<i>Parasisis</i>	sp.	GU338592	GU338500	GU338534	GU338650		

Family	Genus	Species	16s	18s	28s	COI	H3
	<i>Pityophyphantes</i>	<i>costatus</i> *	AY078666	AY078675		AY078695	
	<i>Pocobletus</i>	<i>sp.</i> *	FJ838689	FJ838713		FJ838665	FJ838759
	<i>Porrhomma</i>	<i>sp.</i>	GU338607	GU338466		GU338584	GU338661
	<i>Prosoponoides</i>	<i>sinensis</i>	GU338606			GU338576	GU338649
	<i>Pseudafroneta</i>	<i>incerta</i> *	FJ838690	FJ838714		FJ838666	FJ838760
	<i>Ryotius</i>	<i>sp.</i>	GU338611			GU338536	
	<i>Sisicottus</i>	<i>montanus</i>	GU338625	GU338497		GU338541	GU338673
	<i>Solenysa</i>	<i>sp.</i>	GU338616			GU338528	GU338667
	<i>Sphecozone</i>	<i>bicolor</i>	GU338622	GU338496		GU338553	GU338671
	<i>Stemonyphantes</i>	<i>lineatus</i> *	FJ838691			FJ838738	
	<i>Stemonyphantes</i>	<i>sibiricus</i> *	FJ838692			FJ838667	FJ838761
	<i>Syedra</i>	<i>oii</i>	GU338615	GU338513		GU338666	FJ838762
	<i>Tapinopa</i>	<i>guttata</i>		GU338511		GU338558	
	<i>Tenuiphantes</i>	<i>ancatus</i>		GU338515		GU338567	
	<i>Tenuiphantes</i>	<i>sp.</i>	GU338612	GU338514		GU338568	GU338646
	<i>Tenuiphantes</i>	<i>tenuis</i> *	FJ838693	FJ838716		FJ838739	FJ838763
	<i>Ummeliata</i>	<i>feminea</i>	GU338594	GU338475		GU338551	GU338652
	<i>Ummeliata</i>	<i>insecticeps</i>		GU338476		GU338552	
	<i>Walckenaeria</i>	<i>clavicornis</i>	GU338596	GU338483		GU338554	
	<i>Walckenaeria</i>	<i>keikoa</i>	GU338636	GU338484		GU338556	GU338695